

- VOLUME G -

IN THE UNITED STATES DISTRICT COURT
IN AND FOR THE DISTRICT OF DELAWARE

MONSANTO COMPANY,	:	CIVIL ACTION
	:	
Plaintiff	:	
	:	
	:	
v.	:	
	:	
	:	
MYCOGEN PLANT SCIENCE, INC.,	:	
AGRIGENETICS, INC. and	:	
NOVARTIS CORPORATION,	:	
	:	
Defendants	:	NO. 96-133 (RRM)

Wilmington, Delaware
Thursday, June 25, 1998
8:55 o'clock, a.m.

BEFORE: HONORABLE RODERICK R. McKELVIE, U.S.D.C.J.

APPEARANCES:

POTTER, ANDERSON & CORROON
BY: WILLIAM J. MARSDEN, JR., ESQ.

-and-

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1 A. This is a DNA sequence document that was printed
 2 from the VACS computer at Monsanto at the time I was
 3 there. The FJPERL is Fred's designation. So this would
 4 be one of Fred's documents. And the-BTOURS would
 5 suggest that this is a Bt sequence.
 6 And inside, you see sequence of DNA and
 7 coding sequence. There is yellow highlighting. That
 8 highlighting was characteristic of the way Fred would
 9 mark sequences that he thought might be problematic for
 10 plant expression.
 11 Q. Do you recall ever observing Dr. Perlak make
 12 markings on a document similar to that?
 13 A. Yes.
 14 Q. And when did you first see Dr. Perlak make markings
 15 on a document similar to PTX-157?
 16 A. Within a short time after my arrival, within a few
 17 weeks after my arrival.
 18 Q. Do you recall how frequently you saw Dr. Perlak
 19 working on the Bt redesign project?
 20 A. This was his main focus when I arrived at Monsanto.
 21 He worked on this regularly. Very often.
 22 Q. I am going to hand to the witness what has been
 23 marked as PTX-180. Have you seen PTX-180 before, Dr.
 24 Hanley-Bowdoin?
 25 A. Yes. This is a manuscript and a table that is

1 Perlak?
 2 A. For a year.
 3 Q. Did he work throughout that time on this sequence,
 4 this Bt sequence?
 5 A. Yes.
 6 Q. Did he throughout that time work on highlighting
 7 and modifying printouts such as this?
 8 A. Most of that went on early after I arrived.
 9 Q. Did he continue doing that over a period of time?
 10 A. Yes.
 11 Q. Did he use a number of different documents like
 12 this?
 13 A. I can't speak to that.
 14 Q. Now, you said that somebody showed you this document
 15 earlier today. Did you recognize it when somebody showed
 16 it to you?
 17 A. Yes, I did.
 18 Q. This specific document?
 19 A. I recognize the way this is coded. I recognize
 20 Fred's designation. I recognize the title. I recognize
 21 how it was generated. And I recognize internal to this
 22 the kinds of documents that Fred was using at the time.
 23 Q. The kinds. What about the specific document?
 24 A. I can't speak to that, no, I can't say for a fact
 25 this particular document.

1 codons that are used in genes that are expressed in
 2 various biological systems. So it would be, for example,
 3 a table that you would use to change the codons used in
 4 a bacterial gene to be optimized for plants.
 5 Q. Was PTX-180 readily available in Monsanto?
 6 A. Yes, it was.
 7 Q. Did you ever observe Dr. Perlak using PTX-180?
 8 A. Yes, I did. We had, in our office that we shared,
 9 it was a U-shaped office, and there was a table that
 10 separated our two desks on which there was a computer
 11 that we shared. And over that computer, Fred had these
 12 kinds of tables posted for him to work from. So, yes, I
 13 have seen this in the office.
 14 Q. When was the first time that you recall seeing Dr.
 15 Perlak using Exhibit PTX-180?
 16 A. It was there when I got there.
 17 Q. And did you see him use it?
 18 A. Yes.
 19 Q. Within what time frame?
 20 A. Shortly after I arrived, within a few weeks, again.
 21 MR. LEE: No further questions.
 22 CROSS-EXAMINATION
 23 BY MR. DRIVAS:
 24 Q. Good afternoon, Doctor.
 25 How long did you share an office with Dr.

1 Q. And can you identify any time or date as to when
 2 the changes were made in a document like this?
 3 A. No, I cannot, not specifically.
 4 MR. DRIVAS: Thank you. No further
 5 questions.
 6 MS. MAHANEY: No questions, your Honor.
 7 MR. LEE: No questions.
 8 MR. LYNCH: May Dr. Hanley-Bowdoin be excused,
 9 your Honor?
 10 THE COURT: Yes, you are excused.
 11 (Witness excused)
 12 ---
 13 MS. KNOLL: Your Honor, Monsanto calls Dr.
 14 Tom Rogers.
 15 ---
 16 ... THOMAS E. ROGERS, having been
 17 duly sworn as a witness, was examined and
 18 testified as follows ...
 19 MS. KNOLL: Ladies and gentlemen, Dr. Rogers
 20 is currently an employee with Monsanto. In 1986 he was
 21 in charge of the DNA synthesis group at Monsanto. And he
 22 is going to tell you also about his connections and
 23 relations with Dr. Perlak during that time period.
 24
 25

DIRECT EXAMINATION

3 BY MS. KNOLL:

4 Q. Dr. Rogers, would you please introduce yourself to
5 the jury?

6 A. I am Thomas Rogers from St. Louis, Missouri.

7 Q. What work do you do at Monsanto today?

8 A. Currently I am involved in Human Integrins
9 research, the Pharmaceutical Division at Monsanto.

10 Q. What was your role at Monsanto during the
11 1986/1987 time period?

12 A. At that time, I was the leader of a nucleic acid
13 chemistry crop, within Corporate Research at Monsanto.
14 We had responsibility for all the oligonucleotide
15 synthesis within the company.

16 Q. What did the molecular biology research groups do
17 at Monsanto during that time frame?

18 A. Our function was to provide DNA for all the --
19 anybody who would request DNA for the research purposes
20 at Monsanto.

21 Q. And during that time frame did you come to know a
22 scientist by the name of Fred Perlak?

23 A. Yes, I did.

24 Q. Did you also know Dr. David Fischhoff?

25 A. Yes, I did.

1 Q. How did you come to know those two individuals?

2 A. Well, we would have weekly group meetings,
3 research group meetings, where Fred and Dave would
4 present their research, and we would have a -- usually
5 every month, would have a get-together at an off-site
6 facility and have wine and cheese and that sort of
7 thing. It was just a social sort of thing, in that
8 regard.

9 Then, of course, in our normal daily contact
10 with researchers, Fred and Dave were working
11 oligonucleotides for their projects. So we would get
12 involved with that on a daily basis.

13 Q. Where was your office relative to Dr. Perlak's
14 during the 1986-'87 time frame?

15 A. Fred's office then was due west of mine, about
16 150 feet, on AA-2 in Chesterfield.

17 Q. During that time did you discuss with Dr. Perlak
18 his work on synthetic Bt genes?

19 A. There were two conversations in particular in that
20 regard. Either -- the first one either in late 1986 or
21 early 1987. And the next one was two or three months
22 after that. The first one involved Fred calling me -- I
23 was down at his office, at any rate. And he discussed
24 the problems he was having at that time with Bt expression
25 in plants. And we discussed strategies. In particular,

1 he discussed his strategies for ways to overcome lack of
2 expression of Bt.

3 Q. Let me stop you there and ask you, what was the
4 strategy that Dr. Perlak discussed with you at that
5 time?

6 A. Well, the problem Fred saw was that they were
7 dealing with plant -- bacterial codons, from the parent
8 Bt gene. And his hypothesis was that if we converted
9 that to plant-based codons that the expression levels
10 would then be enhanced in the plant system.

11 Q. Now, Doctor, you mentioned a second conversation.
12 When was the second conversation that you had a
13 recollection about?

14 A. Well, as I mentioned, it was probably two or three
15 months later. Fred was now going to actually begin
16 ordering these oligonucleotides, and he wanted to know
17 whether or not we had the capacity to handle the kind
18 of level of synthesis he was going to require from us.

19 Q. What did you tell him?

20 A. That -- I can't remember precisely what level he
21 was going to require. But that we would have no
22 problems meeting the level of synthesis that he was going
23 to need.

24

25

1

2 Q. Now, when did this second conversation occur?

3 A. The second conversation would have been sometime
4 in -- I would guess in the middle of 1987. Certainly
5 prior to August 22nd, 1987.

6 Q. And how do you know that?

7 A. We went on vacation. My family and I went on
8 vacation on August 22nd, the Saturday the 22nd to Colorado
9 to attend my -- well, first of all, to spend a week at my
10 in-laws in Colorado Springs and then attend my sister's
11 wedding on September 3rd, 1987.

12 Q. Now, let me show you what has previously been marked
13 as two exhibits, Plaintiff's Exhibits 32 and 72. And if
14 you could, Dr. Rogers, tell us what those documents are,
15 just in general terms?

16 A. The first document labeled 32 I believe is a
17 collection of request forms that we would have received
18 from the DNA Chemistry Group. And they begin -- this
19 particular set begins September 24, 1987 from the name
20 of requester is Fred J. Perlak, and then pretty much
21 consecutively in reverse order to November 2nd 1987.
22 Again, these all appear, in this particular number 32,
23 Fred Perlak's.

24 Q. Now, what I would like to ask you first before we
25 get to the specifics of these documents is how did the

1 process work within Monsanto during the '86/'87 time
 2 frame that a researcher like Dr. Perlak would do to get
 3 a piece of DNA synthesized?
 4 A. At that time, the researcher would get on a
 5 computer program on the VACS computer and type in the
 6 sequence that they wanted, the unique name of the file
 7 sequence, and then description of use, and then that
 8 would be mailed to our group, Debbie Connors would
 9 receive these as mail messages. She would print out
 10 these forms as an automatic transfer of this particular
 11 mail message and a form that has information Fred would
 12 have sent or any requester would have sent, the details
 13 about how the synthesizer was going to be set up and how
 14 it would run and also purification scheme.

15 The second page -- I won't read each one of
 16 these -- would be the sequence as we actually would have
 17 put it in. DNA synthesis machine would have been
 18 double-checked by another colleague before we started
 19 synthesis and then the date and name and that sort of
 20 thing. This actually came off the output of the
 21 synthesizer.

22 And then, at the end of the synthesis, and
 23 would have a table again from the synthesizer of actually
 24 what the synthesizer did. So basically the synthesizer
 25 was the robot and this is detailing what the robot did

1 over that period of time.
 2 Q. Now, at the time the researcher or Dr. Perlak sent
 3 these to you, he would have had to tell you what the
 4 sequence was going to be by the time you received it?
 5 A. Yes. I absolutely would have to know the sequence.
 6 Q. Now, the collection that you have in your hand that
 7 is Exhibit 32, these relate to Dr. Perlak's synthetic Bt
 8 work?

9 A. Well if one goes to the description of use, some
 10 cases it's specific. So for example, the item labeled BtK
 11 181.RAQ which is MNP001-054034. The exhibit, description
 12 of use has, to quote:

13 "Sequence for the region of the BtK
 14 gene starting at the base pair 185,
 15 two changes in the codon used."

16 And we go through several of these, if you
 17 wish to show that, indeed, these were related to BtK
 18 genes.

19 Q. Okay. I think that is good for now. We've got an
 20 example.

21 I just have one last question for you, Dr.
 22 Rogers: At this time in Monsanto in 1986/1987, how
 23 common was it for somebody to resynthesize a gene or
 24 rebuild a gene like the one Dr. Perlak was working on?
 25 A. Well, for me to specifically know whether someone

1 is building a gene is fairly rare. Fred came to me
 2 specifically and said this is what we're going to do and
 3 did we have the capacity to do it, which was pretty novel
 4 at that time. Usually we're kept in the dark a little
 5 bit about actually what the use was going to be.

6 MS. KNOLL: Thank you, Dr. Rogers.

7 THE WITNESS: You're welcome.

8 CROSS-EXAMINATION

9 BY MR. DRIVAS:

10 Q. Good afternoon, Dr. Rogers.

11 A. Good afternoon.

12 Q. Are you a chemist by background?

13 A. I'm an organic chemist by background, yes, sir.

14 Q. Do you also design DNA sequences?

15 A. No. That's not generally my forte, although I've

16 done a fair amount of oligonucleotide research.

17 Q. So the practice researchers at Monsanto would

18 actually give you the sequence they wanted synthesized?

19 A. Yes, sir.

20 Q. And then you would have to check that once you made
 21 it?

22 A. Well, once we -- our procedure was, once it was
 23 entered in the machine, we would double-check to make
 24 sure that the sequence fidelity was correct.

25 Q. Is that critical in DNA chemistry?

1 A. Absolutely.

2 Q. Why is that?

3 A. If there is any change you would make could mutate
 4 or probably would mutate depending where it is in the
 5 sequence the particular gene you are trying to modify
 6 and certainly a researcher wants to do a specific change
 7 and not have to fish out something he didn't intend.

8 Q. So even if you made a mistake in one of these
 9 bases, you could have a disastrous effect?

10 A. Could. Depends where it is on the sequence of
 11 the nucleotide.

12 Q. Okay. Now, could you look at Exhibit 32 --

13 A. Sure.

14 Q. -- that was just handed to you? And if you could
 15 turn to the page marked MNP 001-054030...

16 A. (Witness complied.)

17 Q. 030 would be the last three digits?

18 A. Right.

19 Q. Yes.

20 A. I'll get there. Okay.

21 Q. Now, is this typical of the request that you would
 22 get from the researcher?

23 A. Yes.

24 Q. And do you see this BtK 240.RBQ?

25 A. Yes, sir.